AMENDMENTS TO THE CLAIMS

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1-9 cancelled

10. (New) A process for preparing purified 1,3-substituted imidazolium salts of the formula (I)

$$\begin{bmatrix} R2 & R3 \\ R1 & N & R4 \end{bmatrix}^{+} A_{1/a}^{a-}$$
(I),

where

the radicals R¹, R², R³ and R⁴ are each, independently of one another, a carbon-comprising organic, saturated or unsaturated, acyclic or cyclic, aliphatic, aromatic or araliphatic radical which has from 1 to 30 carbon atoms and may comprise one or more heteroatoms and/or be substituted by one or more functional groups or halogens, where adjacent radicals R¹ and R², R² and R³ or R³ and R⁴ may also be joined to one another and the radicals R² and R³ may each also be, independently of one another, hydrogen, halogen or a functional group;

and

A^{a-} is the partly or fully deprotonated anion of an inorganic or organic protic acid H_aA (III), where a is a positive integer and indicates the charge on the anion,

which comprises reacting a 1,3-substituted imidazolium salt of the formula (II),

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$$\begin{bmatrix} R2 & R3 \\ + & \\ R1 - N & + \\ N - R4 \end{bmatrix} + Y_{1/y}^{y-}$$
(II),

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where the radicals R¹, R², R³ and R⁴ are as defined above and

the anion Y^{y^2} is the partly or fully deprotonated anion of an inorganic or organic protic acid

 H_vY (IV),

where y is a positive integer and indicates the charge on the anion,

with a strong base at from 20 to 250°C while distilling off the 1,3-substituted imidazol-2-ylidene formed, wherein the 1,3-substituted imidazol-2-ylidene which has been distilled off is brought into contact in the gaseous state with the protic acid

H_aA (III)

and/or the 1,3-substituted imidazol-2-ylidene which has been distilled off is passed in the gaseous or condensed state into a receiver comprising the protic acid H_aA (III).

- 11. (New) The process according to claim 10, wherein the radials R¹ and R⁴ are each, independently of one another, methyl, ethyl, 1-propyl, 2-propyl, 1-butyl, 1-pentyl, 1-hexyl, 1-heptyl, 1-octyl, 1-nonyl, 1-decyl, 1-dodecyl, 1-tetradecyl, 1-hexadecyl, 1-octadecyl, 1-(2-ethyl)hexyl, benzyl, 3-phenylpropyl, 6-hydroxyhexyl or phenyl and the radicals R² and R³ are each, independently of one another, hydrogen, methyl, ethyl, n-propyl, 2-propyl, 1-butyl, 1-hexyl, 6-hydroxyhexyl, phenyl or chlorine.
- 12. (New) The process according to claim 10, wherein the anion A^{a-} is

fluoride; hexafluorophosphate; hexafluoroarsenate; hexafluoroantimonate; trifluoroarsenate; nitrite; nitrate; sulfate; hydrogensulfate; carbonate; hydrogencarbonate; phosphate; hydrogenphosphate; dihydrogenphosphate; vinyl phosphonate; dicyanamide; bis(pentafluoroethyl)phosphinate; tris(pentafluoroethyl)trifluorophosphate; tris(heptafluoropropyl)trifluorophosphate; bis[oxalato(2-)]borate; bis[salicylato(2-)]borate; bis[1,2-benzenediolato(2-)O,O']borate; tetracyanoborate; tetracarbonylcobaltate;

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tetrasubstituted borate of the formula (Va) [BR^aR^bR^cR^d], where R^a to R^d are each, independently of one another, fluorine or a carbon-comprising organic, saturated or unsaturated, acyclic or cyclic, aliphatic, aromatic or araliphatic radical which has from 1 to 30 carbon atoms and may comprise one or more heteroatoms and/or be substituted by one or more functional groups or halogens;

organic sulfonate of the formula (Vb) [R^e-SO₃], where R^e is a carbon-comprising organic, saturated or unsaturated, acyclic or cyclic, aliphatic, aromatic or araliphatic radical which has from 1 to 30 carbon atoms and may comprise one or more heteroatoms and/or be substituted by one or more functional groups or halogens;

carboxylate of the formula (Vc)

where R^f is hydrogen or a carbon-comprising organic, saturated or unsaturated, acyclic or cyclic, aliphatic, aromatic or araliphatic radical which has from 1 to 30 carbon atoms and may comprise one or more heteroatoms and/or be substituted by one or more functional groups or halogens;

(fluoroalkyl)fluorophosphates of the formula (Vd)

$$[PF_x(C_yF_{2y+1-z}H_z)_{6-x}]^T$$
, (Vd)

where $1 \le x \le 6$, $1 \le y \le 8$ and $0 \le z \le 2y+1$;

imide of the formula (Ve), (Vf) or (IVg)

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$$[R^{g}-SO_{2}-N-SO_{2}-R^{h}]^{-}$$
 (Ve),
 $[R^{i}-SO_{2}-N-CO-R^{j}]^{-}$ (Vf) or
 $[R^{k}-CO-N-CO-R^{l}]^{-}$ (IVg),

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where R^g to R¹ are each, independently of one another, hydrogen or a carbon-comprising organic, saturated or unsaturated, acyclic or cyclic, aliphatic, aromatic or araliphatic radical which has from 1 to 30 carbon atoms and may comprise one or more heteroatoms and/or be substituted by one or more functional groups or halogens; methide of the formula (Vh)

$$\begin{bmatrix} SO_2-R^m \\ | \\ C \\ SO_2-R^o \end{bmatrix}$$
(Vh),

where R^m to R^o are each, independently of one another, hydrogen or a carbon-comprising organic, saturated or unsaturated, acyclic or cyclic, aliphatic, aromatic or araliphatic radical which has from 1 to 30 carbon atoms and may comprise one or more heteroatoms and/or be substituted by one or more functional groups or halogens;

organic sulfate of the formula (Vi) [R^pO-SO₃], where R^p is a carbon-comprising organic, saturated or unsaturated, acyclic or cyclic, aliphatic, aromatic or araliphatic radical which has from 1 to 30 carbon atoms and may comprise one or more heteroatoms and/or be substituted by one or more functional groups or halogens; or

halometalate of the formula (Vj) $[M_qHal_r]^{s-}$, where M is a metal and Hal is fluorine, chlorine, bromine or iodine, q and r are positive integers and indicate the stoichiometry of the complex and s is a positive integer and indicates the charge on the complex.

13. (New) The process according to claim 12, wherein the radials R¹ and R⁴ are each, independently of one another, methyl, ethyl, 1-propyl, 2-propyl, 1-butyl, 1-pentyl, 1-hexyl, 1-

heptyl, 1-octyl, 1-nonyl, 1-decyl, 1-dodecyl, 1-tetradecyl, 1-hexadecyl, 1-octadecyl, 1-(2-ethyl)hexyl, benzyl, 3-phenylpropyl, 6-hydroxyhexyl or phenyl and the radicals R² and R³ are each, independently of one another, hydrogen, methyl, ethyl, n-propyl, 2-propyl, 1-butyl, 1-hexyl, 6-hydroxyhexyl, phenyl or chlorine.

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- 14. (New) The process according to claim 10, wherein the anion A^{a-} is tetrafluoroborate, hexafluorophosphate, trifluoromethanesulfonate, methanesulfonate, formate, acetate, mandelate, nitrate, nitrite, trifluoroacetate, sulfate, hydrogensulfate, methyl sulfate, ethyl sulfate, propyl sulfate, butyl sulfate, pentyl sulfate, hexyl sulfate, heptyl sulfate, octyl sulfate, phosphate, dihydrogenphosphate, hydrogenphosphate, propionate, tetrachloroaluminate, Al₂Cl₇, chlorozincate, chloroferrate, bis(trifluoromethylsulfonyl)imide, bis(pentafluoroethylsulfonyl)imide, tris(trifluoromethylsulfonyl)methide, bis(pentafluoroethylsulfonyl)methide, p-toluenesulfonate, bis[salicylato(2-)]borate, tetracarbonylcobaltate, dimethylene glycol monomethyl ether sulfate, octyl sulfate, oleate, stearate, acrylate, methacrylate, maleate, hydrogencitrate, vinyl phosphonate, bis(pentafluoroethyl)phosphinate, bis[oxalato(2-)]borate, bis[1,2-benzenediolato(2-)O,O']borate, dicyanamide, tris(pentafluoroethyl)trifluorophosphate, tris(heptafluoropropyl)trifluorophosphate, tetracyanoborate or chlorocobaltate.
- 15. (New) The process according to claim 13, wherein the anion A^{a-} is tetrafluoroborate, hexafluorophosphate, trifluoromethanesulfonate, methanesulfonate, formate, acetate, mandelate, nitrate, nitrite, trifluoroacetate, sulfate, hydrogensulfate, methyl sulfate, ethyl sulfate, propyl sulfate, butyl sulfate, pentyl sulfate, hexyl sulfate, heptyl sulfate, octyl sulfate, phosphate, dihydrogenphosphate, hydrogenphosphate, propionate, tetrachloroaluminate, Al₂Cl₇, chlorozincate, chloroferrate, bis(trifluoromethylsulfonyl)imide, bis(pentafluoroethylsulfonyl)imide, tris(trifluoromethylsulfonyl)methide, bis(pentafluoroethylsulfonyl)methide, p-toluenesulfonate, bis[salicylato(2-)]borate, tetracarbonylcobaltate, dimethylene glycol monomethyl ether sulfate, octyl sulfate, oleate, stearate, acrylate, methacrylate, maleate, hydrogencitrate, vinyl phosphonate, bis(pentafluoroethyl)phosphinate, bis[oxalato(2-)]borate, bis[1,2-benzenediolato(2-)O,O']borate, dicyanamide, tris(pentafluoroethyl)trifluorophosphate, tris(heptafluoropropyl)trifluorophosphate, 439479

tetracyanoborate or chlorocobaltate.

16. (New) The process according to claim 10, wherein the anion Y^{y-} is chloride, bromide, methanesulfonate, hydrogencarbonate, carbonate, hydrogensulfate, diethylphosphate, tosylate or methyl sulfate.

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- 17. (New) The process according to claim 15, wherein the anion Y^y is chloride, bromide, methanesulfonate, hydrogencarbonate, carbonate, hydrogensulfate, diethylphosphate, tosylate or methyl sulfate.
- 18. (New) The process according to claim 10, wherein the 1,3-substituted imidazol-2-ylidene which has been distilled off is brought into contact in the gaseous state with gaseous protic acid H_aA (III) and the condensed, purified 1,3-substituted imidazolium salt (I) is isolated.
- 19. (New) The process according to claim 17, wherein the 1,3-substituted imidazol-2-ylidene which has been distilled off is passed in the gaseous state into a receiver comprising the protic acid H_aA (III) and the purified 1,3-substituted imidazolium salt (I) is isolated therefrom.
- 20. (New) The process according to claim 10, wherein the 1,3-substituted imidazol-2-ylidene which has been distilled off is condensed in a condenser, passed in the condensed state into a distillation receiver comprising the protic acid H_aA (III) and the purified 1,3-substituted imidazolium salt (I) is isolated therefrom.
- 21. (New) The process according to claim 19, wherein the 1,3-substituted imidazol-2-ylidene which has been distilled off is condensed in a condenser, passed in the condensed state into a distillation receiver comprising the protic acid H_aA (III) and the purified 1,3-substituted imidazolium salt (I) is isolated therefrom.
- 22. (New) The process according to claim 10, wherein the distillation is carried out at a pressure of from 0.0001 to 0.15 MPa abs.

23. (New) The process according to claim 21, wherein the distillation is carried out at a pressure of from 0.0001 to 0.15 MPa abs.

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